

What is claimed is:

1. A method of treating a patient having melanoma which has been surgically removed, which comprises administering to such a patient a therapeutically effective dose of pegylated interferon alpha for a time period sufficient to increase progression-free survival time.
2. The method of claim 1 wherein the pegylated interferon is pegylated interferon alpha-2a or pegylated interferon alpha-2b.
3. The method of claim 2 wherein the patient is a treatment-naive patient.
4. The method of claim 3 wherein the treatment-naive patient is one having newly diagnosed melanoma.
5. The method of claim 1 wherein the patient is treatment-experienced patient.
6. The method of claim 5 wherein the treatment experienced patient is intolerant to interferon alpha or resistant to interferon alpha.
7. The method of claim 1 wherein the time period is at least about 24 months.
8. The method of claim 1 wherein the pegylated interferon alpha administered after surgical excision of the primary melanoma.
9. The method of claim 1 wherein the pegylated interferon alpha is pegylated interferon alpha-2b and the effective amount is in the range of about 3.0 micrograms/kg to 9.0 micrograms/kg administered once a week.

10. The method of claim 1 wherein the pegylated interferon alpha is pegylated alpha-2a and the effective amount is in the range of about 200 microgram to 250 administered once a week.
11. A method of treating a patient having cutaneous melanoma which has been surgically removed which comprises administering to said patient an effective amount of pegylated interferon-alpha once a week for a time period sufficient to increase progression-free survival time.
12. The method of claim 11 wherein the pegylated interferon alpha is pegylated interferon alpha-2b and the effective amount is in the range of about 3.0 micrograms/kg to 9.0 micrograms/kg administered once a week.
13. The method of claim 11 wherein the pegylated interferon alpha is pegylated alpha-2a and the effective amount is in the range of about 200 microgram to 250 administered once a week.
14. The method of claim 11 wherein the time period is at least about 100 weeks.
15. A method of treating a patient having cutaneous melanoma which comprises administering to such a patient about 3.0 micrograms/kg to about 9.0 micrograms/kg of pegylated interferon alpha-2b once a week for a time period sufficient to increase the progression-free survival time.
16. The method of claim 15 wherein the time period is about 100 weeks.
17. The method of claim 15 wherein about 4.5 to about 6.5 micrograms/kg of pegylated interferon alpha-2b is administered once a week.

18. A method comprising the step of marketing a therapeutically effective dose of pegylated interferon alpha for administration to a patient with melanoma within about 60 days of surgery in a protocol extending for a time period of at least about 100 weeks.
19. The method of claim 18 wherein the pegylated interferon alpha is pegylated interferon alpha-2b and the effective amount is in the range of about 3.0 micrograms/kg to 9.0 micrograms/kg administered once a week.
20. The method of claim 18 wherein the pegylated interferon alpha is pegylated alpha-2a and the effective amount is in the range of about 200 microgram to 250 administered once a week.

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